

(12) INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(19) World Intellectual Property Organization  
International Bureau



(43) International Publication Date  
25 May 2001 (25.05.2001)

PCT

(10) International Publication Number  
**WO 01/36384 A1**

(51) International Patent Classification<sup>7</sup>: C07D 207/335,  
A61K 31/40

Even-Yeouda (IL). LIFSHITZ, Revital [IL/IL]; 12a/8  
Kibush Haavoda, 46322 Herzilia (IL).

(21) International Application Number: PCT/US00/31555

(74) Agents: BRAINARD, Charles, R. et al.; Kenyon &  
Kenyon, One Broadway, New York, NY 10004 (US).

(22) International Filing Date:

16 November 2000 (16.11.2000)

(25) Filing Language:

English

(26) Publication Language:

English

(30) Priority Data:

60/166,153 17 November 1999 (17.11.1999) US

(71) Applicant (for all designated States except BB, US):  
TEVA PHARMACEUTICAL INDUSTRIES LTD.  
[IL/IL]; Basel Street 5, P.O. Box 3190, 49131 Petah Tiqva  
(IL).

(81) Designated States (national): AE, AG, AL, AM, AT, AU,  
AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN, CR, CU, CZ,  
DE, DK, DM, DZ, EE, ES, FI, GB, GD, GE, GH, GM, HR,  
HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR,  
LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ,  
NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM,  
TR, TT, TZ, UA, UG, US, UZ, VN, YU, ZA, ZW.

(84) Designated States (regional): ARIPO patent (GH, GM,  
KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW), Eurasian  
patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European  
patent (AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE,  
IT, LU, MC, NL, PT, SE, TR), OAPI patent (BF, BJ, CF,  
CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG).

(71) Applicant (for BB only): TEVA PHARMACEUTICALS  
USA, INC. [US/US]; 1090 Horsham Road, P.O. Box 1090,  
North Wales, PA 19454-1090 (US).

**Published:**

- With international search report.
- Before the expiration of the time limit for amending the  
claims and to be republished in the event of receipt of  
amendments.

(72) Inventors; and

(75) Inventors/Applicants (for US only): AYALON, Ari  
[IL/IL]; HaAri Street 8, Haifa (IL). LEVINGER, Michael  
[IL/IL]; Rehov Eitanim, P.O. Box 207, Nofit (IL). ROYT-  
BLAT, Sofia [IL/IL]; Netiv Chen 6, Haifa (IL). NIDDAM,  
Valerie [IL/IL]; 9 Keren Hayessod, P.O. Box 1343, 40500

For two-letter codes and other abbreviations, refer to the "Guid-  
ance Notes on Codes and Abbreviations" appearing at the begin-  
ning of each regular issue of the PCT Gazette.

WO 01/36384 A1

(54) Title: POLYMORPHIC FORM OF ATORVASTATIN CALCIUM

(57) Abstract: The present invention relates to atorvastatin calcium, a useful agent for lowering serum cholesterol levels. New atorvastatin calcium Form V, processes for preparing the new form, and pharmaceutical compositions and dosage forms containing the new form are disclosed.

## POLYMORPHIC FORM OF ATORVASTATIN CALCIUM

FIELD OF THE INVENTION

The present invention relates to a novel crystalline form of atorvastatin calcium, to  
5 processes for preparing it and to pharmaceutical compositions and dosages containing it.

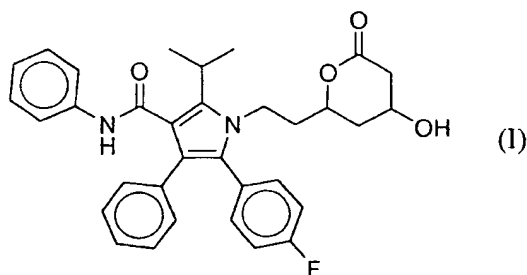
BACKGROUND OF THE INVENTION

Atorvastatin is a member of the class of drugs called statins. Statin drugs are  
currently the most therapeutically effective drugs available for reducing low density  
10 lipoprotein (LDL) particle concentration in the blood stream of patients at risk for  
cardiovascular disease. A high level of LDL in the bloodstream has been linked to the  
formation of coronary lesions which obstruct the flow of blood and can rupture and  
promote thrombosis. Goodman and Gilman. *The Pharmacological Basis of Therapeutics*  
879 (9th ed. 1996). Reducing plasma LDL levels has been shown to reduce the risk of  
15 clinical events in patients with cardiovascular disease and patients who are free of  
cardiovascular disease but who have hypercholesterolemia. Scandinavian Simvastatin  
Survival Study Group, 1994; Lipid Research Clinics Program, 1984a, 1984b.

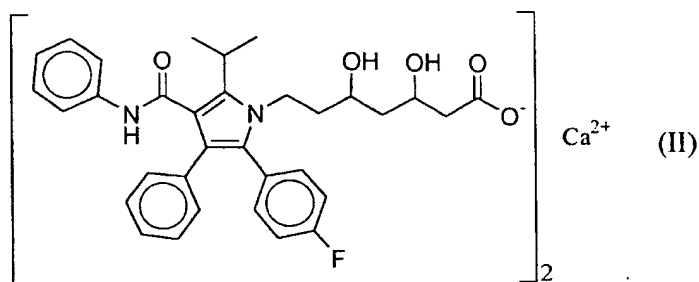
The mechanism of action of statin drugs has been elucidated in some detail. They  
interfere with the synthesis of cholesterol and other sterols in the liver by competitively  
20 inhibiting the 3-hydroxy-3-methyl-glutaryl-coenzyme A reductase enzyme ("HMG-CoA  
reductase"). HMG-CoA reductase catalyzes the conversion HMG to mevalonate, which

is the rate determining step in the biosynthesis of cholesterol, and so, its inhibition leads to a reduction in the concentration of cholesterol in the liver. Very low density lipoprotein (VLDL) is the biological vehicle for transporting cholesterol and triglycerides from the liver to peripheral cells. VLDL is catabolized in the peripheral cells which releases fatty acids which may be stored in adipocytes or oxidized by muscle. The VLDL is converted to intermediate density lipoprotein (IDL), which is either removed by an LDL receptor, or is converted to LDL. Decreased production of cholesterol leads to an increase in the number of LDL receptors and corresponding reduction in the production of LDL particles by metabolism of IDL.

Atorvastatin is the common chemical name of  $[R-(R^*, R^*)]$ -2-(4-fluorophenyl)- $\beta,\delta$ -dihydroxy-5-(1-methylethyl)-3-phenyl-4-[(phenylamino)carbonyl]-1H-pyrrole-1-heptanoic acid. The free acid is prone to lactonization. The molecular structure of the lactone is represented by formula (I).



Atorvastatin is marketed as the hemi calcium salt-trihydrate under the name LIPITOR by Warner-Lambert Co.



Atorvastatin was first disclosed to the public and claimed in U.S. Patent No. 4,681,893. The hemi calcium salt depicted in formula (II) (hereafter "atorvastatin calcium") is disclosed in U.S. Patent No. 5,273,995. This patent teaches that the calcium salt is obtained by crystallization from a brine solution resulting from the transposition of the sodium salt with  $\text{CaCl}_2$  and further purified by recrystallization from a 5:3 mixture of ethyl acetate and hexane. Both of these U.S. patents are hereby incorporated by reference.

The present invention includes a new crystal form of atorvastatin calcium in both hydrate and anhydrate states. Polymorphism is the property of some molecules and molecular complexes to assume more than one crystalline or amorphous form in the solid state. A single molecule, like the atorvastatin in formula (I) or the salt complex of formula (II), may give rise to a variety of solids having distinct physical properties like solubility, X-ray diffraction pattern and solid state  $^{13}\text{C}$  NMR spectrum. The differences in the physical properties of polymorphs result from the orientation and intermolecular interactions of adjacent molecules (complexes) in the bulk solid. Accordingly, polymorphs are distinct solids sharing the same molecular formula, which may be thought

of as analogous to a unit cell in metallurgy, yet having distinct advantageous and/or disadvantageous physical properties compared to other forms in the polymorph family. One of the most important physical properties of pharmaceutical polymorphs is their solubility in aqueous solution, particularly their solubility in the gastric juices of a patient.

5 For example, where absorption through the gastrointestinal tract is slow, it is often desirable for a drug that is unstable to conditions in the patient's stomach or intestine to dissolve slowly so that it does not accumulate in a deleterious environment. On the other hand, where the effectiveness of a drug correlates with peak bloodstream levels of the drug, a property shared by statin drugs, and provided the drug is rapidly absorbed by the  
10 GI system, then a more rapidly dissolving form is likely to exhibit increased effectiveness over a comparable amount of a more slowly dissolving form.

U.S. Patent No. 5,969,156 discloses three polymorphs of atorvastatin designated Forms I, II, and IV by the inventors of those forms. While the inventors of U.S. Patent No. 5,969,156 claim certain processing and therapeutic advantages of their forms over  
15 amorphous atorvastatin calcium, advantages may yet be realized by other heretofore undiscovered forms of atorvastatin calcium.

#### SUMMARY OF THE INVENTION

The present invention provides new Form V of atorvastatin calcium in both  
20 anhydrate and hydrate states, which possesses the advantage of higher solubility in water than atorvastatin Form I. The present invention further provides a process for preparing

new Form V as well as pharmaceutical compositions and dosages containing the new form.

#### BRIEF DESCRIPTION OF THE DRAWINGS

5 FIG. 1 is an X-ray powder diffractogram of atorvastatin calcium Form V.

FIG. 2 is a solid state  $^{13}\text{C}$  NMR spectrum of atorvastatin calcium Form V.

#### DETAILED DESCRIPTION OF THE INVENTION

The new crystalline form of atorvastatin calcium Form V is well distinguished  
10 from the crystal modifications obtained by carrying out the procedures described in U.S. Patents Nos. 5,273,995 and 5,969,156 using X-ray powder diffraction and solid-state  $^{13}\text{C}$  nuclear magnetic resonance techniques.

The X-ray powder diffractogram of Form V (Fig.1) has two medium peaks at  
5.3 $\pm$ 0.2 and 8.3 $\pm$ 0.2 degrees 2 $\theta$  and one large peak in the range 18-23 degrees 2 $\theta$  with a  
15 maximum at about 18.3 $\pm$ 0.2 degrees two-theta. This X-Ray pattern is well distinguished from that of known Forms I, II, III and IV and also is well distinguished from the X-Ray pattern of amorphous atorvastatin calcium which is characterized by two broad humps in the ranges 8-14 degrees 2 $\theta$  and 15-26 degrees 2 $\theta$ . The X-ray powder diffractogram of  
Fig. 1 was obtained by methods known in the art using a Philips X-ray powder  
20 diffractometer with a curved graphite monochromator using goniometer model 1050/70. Copper radiation of  $\lambda = 1.5418 \text{ \AA}$  was used. Measurement range: 3-30 degrees 2 $\theta$ .

The solid-state  $^{13}\text{C}$  NMR spectrum of Form V is characterized by the following chemical shifts:

	$\delta$ (ppm)
	21.9
5	25.9
	40.4
	41.8
	42.3
	63-73 (two broad peaks)
10	115.6
	118.9
	122.5
	128.7 (strong)
	135.1
15	161.0
	167.1
	176-186 (broad peak)

This solid-state  $^{13}\text{C}$  NMR spectrum (Fig. 2) is well distinguished from those of known Forms I, II, III and IV, and also is distinguished from that of the amorphous form which displays a different pattern with shifts significantly different from that of Form V at 21.0 ppm, 26.4 ppm, one broad peak in the range 60-75 ppm with a maximum at 69.7 ppm and 138.8 ppm. The spectrum of Fig. 2 was obtained on a Bruker DMX-500 digital F NMR spectrometer operating at 125.76 MHz. The instrument was equipped with a BL-4 cpmas probehead and a high resolution/high performance (HPHP)  $^1\text{H}$  for solids. The magic angle and proton decoupling efficiency were optimized before acquisition. The sample was spun at 5.0 kHz spin rate on 4 mm zirconia rotors.

Atorvastatin calcium Form V may contain up to 12% water, which corresponds to the stoichiometric value of 9 water molecules per molecule of atorvastatin calcium. Thus, atorvastatin calcium Form V can be in various states of hydration, between 0 and 9 moles of water.

5           The present invention further provides a process for the preparation of atorvastatin calcium Form V. The process comprises the steps of dissolving a salt of atorvastatin in a solvent to form an atorvastatin salt solution, optionally removing impurities from the atorvastatin salt solution, contacting the atorvastatin salt solution with a calcium salt and isolating atorvastatin calcium in new Form V.

10           The atorvastatin salt of the present invention includes alkali metal salts, e.g. lithium, sodium, and potassium salts; alkaline-earth metal salts such as magnesium salts; as well as ammonium and alkyl, aryl or alkaryl ammonium salts. The preferred atorvastatin salts are alkali metal salts; most preferred is the sodium salt.

Any solvent capable of dissolving the atorvastatin salt and from which atorvastatin  
15 calcium Form V may be isolated is a suitable solvent of the invention. The choice of solvent will therefore depend upon the selection of the atorvastatin salt and the calcium salt. The solvent should be selected from those in which the atorvastatin salt and calcium salt are at least sparingly soluble. By sparingly soluble is meant not substantially less  
soluble than 0.02 g/ml at 50-60°C for the atorvastatin salt and not substantially less  
20 soluble than 0.0002 M at 10-15°C for the calcium salt.



Suitable solvents include but are not limited to hydroxylic solvents like water, alcohols and mixtures thereof, including hydroxylic solvents and hydroxylic solvent mixtures which have been made either acidic or basic by addition of a mineral acid or base. Preferred solvents are water, methanol, ethanol and mixtures thereof.

- 5           The calcium salt of the present invention includes organic and inorganic salts of calcium which are capable of dissociating into  $\text{Ca}^{2+}$  and an anionic component when added to the atorvastatin salt solution. Among the organic salts that may be used are carboxylates and sulfonates. Among the carboxylates are lower alkyl carboxylates like acetate, propionate, butyrate and tartrate and aryl carboxylates like benzoate and
- 10           phthalate as well as higher alkyl carboxylates like stearate, dodecanoate and the like. Also included are calcium ascorbate and succinate. Among the sulfonates that may be used are lower alkyl and aryl sulfonates like calcium methane sulfonate, calcium benzene sulfonate and calcium *p*-toluene sulfonate. The preferred organic calcium salts are lower carboxylate salts, the most preferred organic calcium salt is calcium acetate.
- 15           Depending upon solubility, inorganic salts which may be used include halide salts such as  $\text{CaCl}_2$ ,  $\text{CaF}_2$ ,  $\text{CaBr}_2$  and  $\text{CaI}_2$ , as well as calcium borate ( $\text{B}_4\text{CaO}_7$ ), calcium tetrafluoroborate ( $\text{CaBF}_4$ ), calcium carbonate ( $\text{CaCO}_3$ ), monobasic calcium phosphate ( $\text{Ca}(\text{H}_2\text{PO}_4)_2$ ), dibasic calcium phosphate ( $\text{CaHPO}_4$ ) and tribasic calcium phosphate ( $\text{Ca}(\text{PO}_4)_2$ ), calcium sulfate ( $\text{CaSO}_4$ ) and calcium hydroxide ( $\text{Ca}(\text{OH})_2$ ), and hydrates
- 20           thereof.

Whether organic or inorganic, the calcium salt is preferably added in an amount that provides one half mole of  $\text{Ca}^{2+}$  per mole of atorvastatin in the atorvastatin salt solution. For example, if the atorvastatin salt is atorvastatin sodium ( $\text{atorvastatin}^- \text{Na}^+$ ), then about one half mole of calcium salt per mole of the atorvastatin salt is appropriate. If  
5 the atorvastatin salt is atorvastatin magnesium ( $[\text{atorvastatin}^-]_2 \text{Mg}^{2+}$ ), then about one mole of calcium salt per mole of atorvastatin salt is appropriate. Otherwise, mixed salts containing atorvastatin may form.

The calcium salt may be contacted with the atorvastatin salt solution by adding the calcium salt in substantially pure form, *i.e.* either as a solid or, if liquid, as a neat liquid, to  
10 the atorvastatin salt solution or, preferably, by first forming a calcium salt solution and then contacting the atorvastatin salt solution and calcium salt solution. It is most preferred to contact the calcium salt and the atorvastatin salt solution by first dissolving the calcium salt in a solvent and then adding the calcium salt solution to the atorvastatin salt solution slowly. Suitable calcium salt solvents are solvents previously mentioned as being suitable  
15 solvents for the atorvastatin salt, provided the calcium salt is at least sparingly soluble in the particular solvent.

In a particularly preferred embodiment, wherein the atorvastatin salt is an atorvastatin alkali metal salt and the atorvastatin salt solvent is a 1:2 methanol:water mixture, the preferred calcium salt is calcium acetate and the preferred calcium salt solvent  
20 is water. When the calcium salt solvent is water, it is preferably used in an amount that

provides about a 20 to 30 millimolar solution of the calcium salt, more preferably about a 25 millimolar solution.

In addition, the atorvastatin and calcium salts are preferably combined at elevated temperature and at concentrations disclosed above and in the examples, which follow, in order that crystallization of Form V may be induced by cooling of the so-formed atorvastatin calcium solution. The elevated temperature is preferably above 40°C and below 80°C, more preferably above 50°C and below 70°C and most preferably about 60°C. One skilled in the art will appreciate that by adjusting temperature and concentration, the yield of atorvastatin calcium Form V may be optimized. Crystallization of atorvastatin calcium Form V may also be induced by addition of a seed crystal of atorvastatin calcium, preferably Form V although other forms also may be used.

Once crystals of atorvastatin Form V have crystallized, either spontaneously, upon cooling, upon seeding or by another inducement, the crystals may be isolated by filtration or other conventional means known to the art. The isolated crystals may also be dried by conventional means.

It has also been found that atorvastatin calcium can be crystallized in Form V by dissolving atorvastatin calcium in THF or alcohols like methanol or ethanol, and subsequently adding water as an antisolvent.

A further aspect of the present invention is a pharmaceutical composition and dosage form containing the novel form of atorvastatin calcium.

The compositions of the invention include powders, granulates, aggregates and other solid compositions comprising novel Form V of atorvastatin calcium. In addition, Form V solid compositions that are contemplated by the present invention may further included diluents, such as cellulose-derived materials like powdered cellulose, microcrystalline cellulose, microfine cellulose, methyl cellulose, ethyl cellulose, hydroxyethyl cellulose, hydroxypropyl cellulose, hydroxypropylmethyl cellulose, carboxymethyl cellulose salts and other substituted and unsubstituted celluloses; starch; pregelatinized starch; inorganic diluents like calcium carbonate and calcium diphosphate and other diluents known to the pharmaceutical industry. Yet other suitable diluents include waxes, sugars and sugar alcohols like mannitol and sorbitol, acrylate polymers and copolymers, as well as pectin, dextrin and gelatin.

Further excipients that are within the contemplation of the present invention include binders, such as acacia gum, pregelatinized starch, sodium alginate, glucose and other binders used in wet and dry granulation and direct compression tableting processes. Excipients that may also be present in a solid composition of Form V atorvastatin calcium further include disintegrants like sodium starch glycolate, croscopovidone, low-substituted hydroxypropyl cellulose and others. In addition, excipients may include tableting lubricants like magnesium and calcium stearate and sodium stearyl fumarate; flavorings; sweeteners; preservatives; pharmaceutically acceptable dyes and glidants such as silicon dioxide.

The dosages include dosages suitable for oral, buccal, rectal, parenteral (including subcutaneous, intramuscular, and intravenous), inhalant and ophthalmic administration. Although the most suitable route in any given case will depend on the nature and severity of the condition being treated, the most preferred route of the present invention is oral. The  
5 Dosages may be conveniently presented in unit dosage form and prepared by any of the methods well-known in the art of pharmacy.

Dosage forms include solid dosage forms, like tablets, powders, capsules, suppositories, sachets, troches and lozenges as well as liquid suspensions and elixirs. While the description is not intended to be limiting, the invention is also not intended to  
10 pertain to true solutions of atorvastatin calcium whereupon the properties that distinguish the solid forms of atorvastatin calcium are lost. However, the use of the novel forms to prepare such solutions (e.g. so as to deliver, in addition to atorvastatin, a solvate to said solution in a certain ratio with a solvate) is considered to be within the contemplated invention.

15 Capsule dosages, of course, will contain the solid composition within a capsule which may be made of gelatin or other conventional encapsulating material. Tablets and powders may be coated. Tablets and powders may be coated with an enteric coating. The enteric coated powder forms may have coatings comprising phthalic acid cellulose acetate, hydroxypropylmethyl-cellulose phthalate, polyvinyl alcohol phthalate,  
20 carboxymethylethylcellulose, a copolymer of styrene and maleic acid, a copolymer of methacrylic acid and methyl methacrylate, and like materials, and if desired, they may be

employed with suitable plasticizers and/or extending agents. A coated tablet may have a coating on the surface of the tablet or may be a tablet comprising a powder or granules with an enteric-coating.

Preferred unit dosages of the pharmaceutical compositions of this invention typically contain from 0.5 to 100 mg of the novel atorvastatin calcium Form V, or mixtures thereof with other forms of atorvastatin calcium. More usually, the combined weight of the atorvastatin calcium forms of a unit dosage are from 2.5 mg. to 80 mg.

Having thus described the various aspects of the present invention, the following examples are provided to illustrate specific embodiments of the present invention. They are not intended to be limiting in any way.

## EXAMPLES

### EXAMPLE 1

The sodium salt of atorvastatin (52.2 g) was dissolved in methanol (510 ml) and then diluted with water (1 L). The resulting solution was transferred to a separatory funnel containing 1:1 ethyl acetate/hexane (1 L). The phases were mixed by bubbling nitrogen gas through the separatory funnel. Upon cessation of nitrogen flow, the phases separated and the upper, organic, phase was removed. The lower, aqueous, phase was washed with 1:1 ethyl acetate/hexane (1 L) and then transferred to a round bottom flask. Active charcoal (10.2 g) was added. The flask was heated to 50 °C and the solution was stirred for two hours. The activated charcoal was then removed by filtration through celite, the

charcoal and celite being rinsed with methanol (1540 ml), and the rinsate and filtrate then being combined into one atorvastatin sodium salt solution.

The quantity of atorvastatin obtained by purification was determined by calibrated HPLC analysis of the purified atorvastatin sodium salt solution. Based on this analysis, a  
5 quantity of calcium acetate (8.38 g, 0.5 eq.) was dissolved in water (1.9 L) and heated to 60 °C. The atorvastatin sodium salt solution was heated to 63 °C and the solutions were combined by slow addition of the calcium acetate solution to the atorvastatin sodium salt solution. Upon completing the addition, the mixture was cooled. Crystallization of Form V began to occur at a temperature of 43 °C and cooling was continued until the flask  
10 temperature reached 13 °C.

The crystals were isolated by slow vacuum filtration and then dried over anhydrous silica for 5 days to yield atorvastatin calcium salt Form V.

## EXAMPLE 2

15 Atorvastatin calcium (10 g) was dissolved in methanol (400 ml) at room temperature. Water (300 ml) was added slowly to the methanolic solution with stirring and the resulting solution was heated to 60°C. The solution was then cooled to between 10 and 15 °C within 3 h. Precipitation started at about 40°C. The thick slurry was then dried at 50°C under reduced pressure for 48 h to yield atorvastatin calcium Form V.

20

## EXAMPLE 3

Atorvastatin calcium (5 g) was dissolved in methanol (100 ml) at room temperature. To this methanolic solution, water (100 ml) was added while stirring. Precipitation occurred instantly and after cooling the slurry to 15°C the precipitate was  
5 filtered and dried at 50°C under reduced pressure for 48 h to yield atorvastatin calcium Form V.

## EXAMPLE 4

Atorvastatin calcium (5 g) was dissolved in methanol (200 ml). The methanolic  
10 solution was placed into a stirred reactor containing water (150 ml) at 45°C. The obtained slurry was cooled to 10°C, filtered and dried at 50°C under reduced pressure for 48 h to yield atorvastatin calcium Form V.

## EXAMPLE 5

15 Atorvastatin calcium (1 g) was dissolved in ethanol (15 ml) after heating. To this ethanolic solution, water (10 ml) was added while stirring. Precipitation occurred instantly. The gel-like precipitate was filtered without vacuum and dried at 50°C under reduced pressure for 24 h to yield atorvastatin calcium Form V.



## EXAMPLE 6

Atorvastatin calcium (1 g) was dissolved in THF (25 ml) at room temperature. To this solution, water (60 ml) was added while stirring. The reaction mixture was stirred for 18 hours at room temperature and the precipitate (gel) was filtered without vacuum and  
5 dried at 50°C under reduced pressure for 24 h to yield atorvastatin calcium Form V.

The invention has been described with reference to its preferred embodiments. From this description, those skilled in the art may appreciate changes that could be made in the invention which do not depart from the scope and spirit of the invention as described above and claimed hereafter.

## CLAIMS

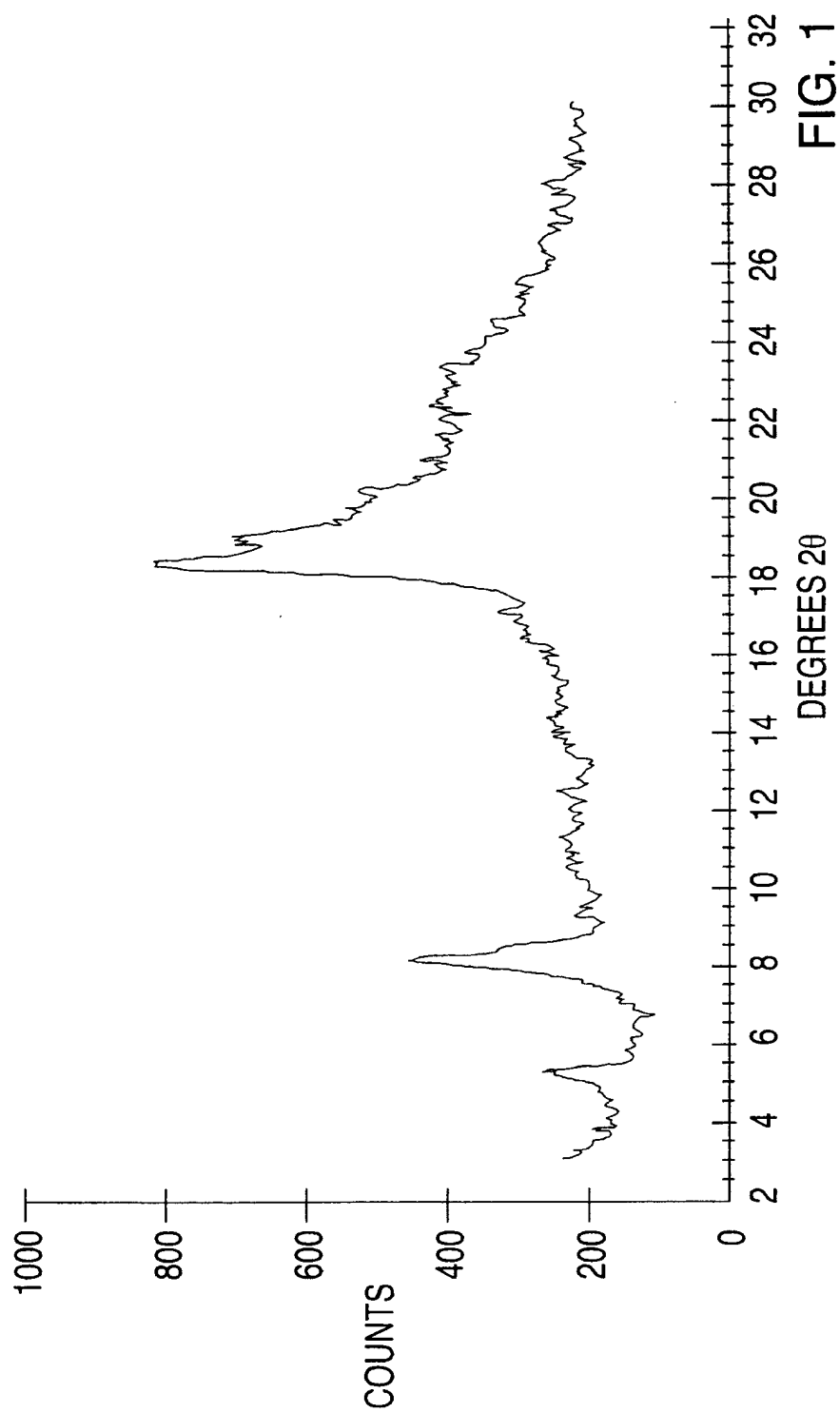
We claim:

1. Atorvastatin calcium Form V or hydrate thereof.
2. The atorvastatin calcium Form V or hydrate thereof of claim 1 having an X-ray powder diffractogram substantially as depicted in Figure 1.
3. The atorvastatin calcium Form V or hydrate thereof of claim 1 having X-ray powder diffraction peaks at about 5.5 and 8.3 degrees at  $2\theta$  and a broad peak at about 18-23 degrees  $2\theta$ .
4. The atorvastatin calcium Form V or hydrate thereof of claim 1 having a solid state  $^{13}\text{C}$  NMR spectrum substantially as depicted in Figure 2.
5. The atorvastatin calcium Form V or hydrate thereof of claim 1 having solid state  $^{13}\text{C}$  NMR signals at about 21.9, 25.9, 118.9, 122.5, 128.7, 161.0 and 167.1 ppm.
6. Atorvastatin calcium Form V containing up to about 9 moles of water per mole of atorvastatin calcium.
7. A process for preparing atorvastatin calcium Form V, or hydrate thereof, comprising the steps of
  - a) dissolving a metal, ammonium or alkylammonium salt of atorvastatin in a solvent to form an atorvastatin salt solution
  - b) contacting the atorvastatin salt solution with a calcium salt, and
  - c) isolating atorvastatin calcium Form V or hydrates thereof.
8. The process of claim 7 wherein the salt of atorvastatin is a metal salt of

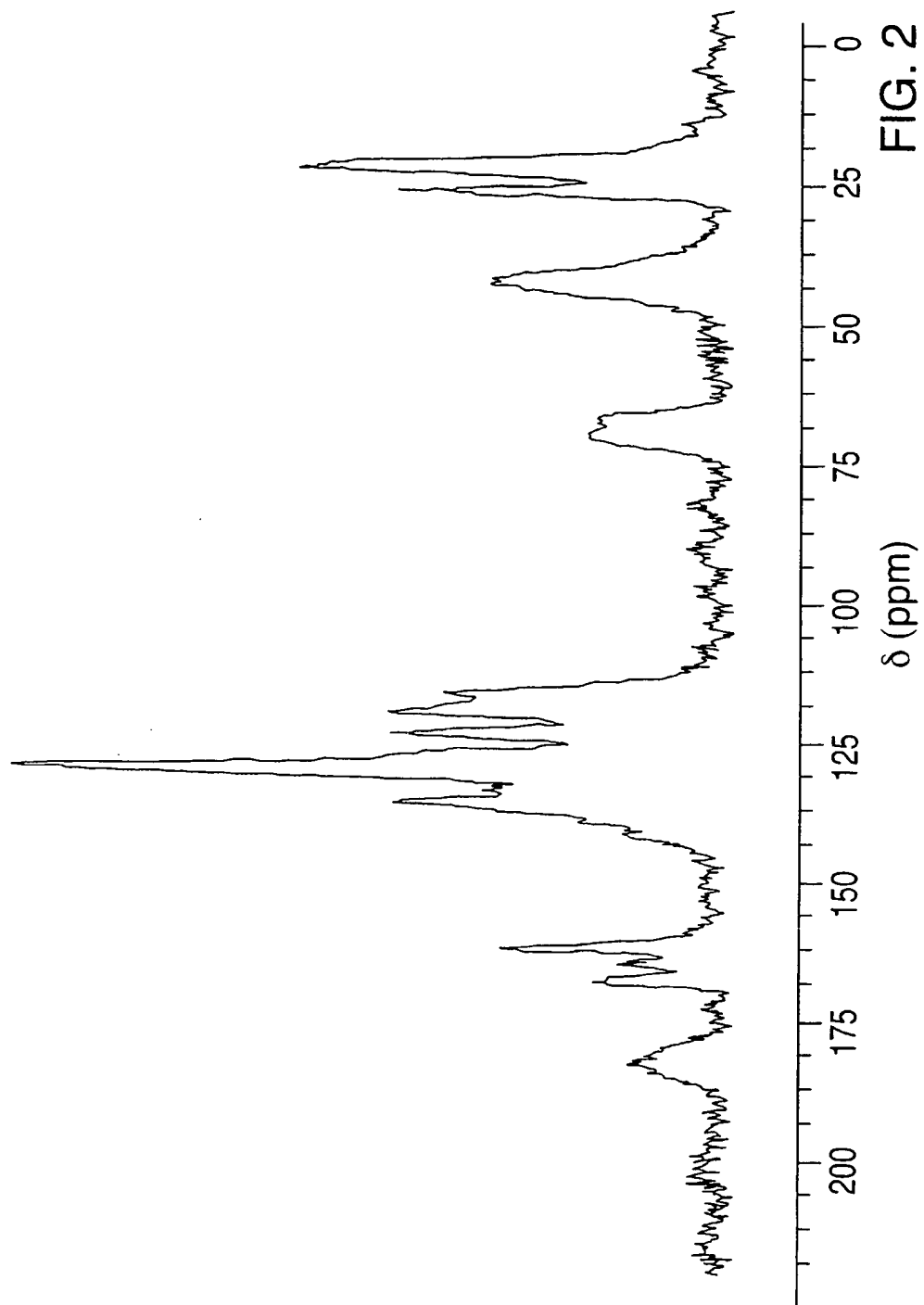
atorvastatin.

9. The process of claim 8 wherein the metal salt of atorvastatin is a sodium salt of atorvastatin.
10. The process of claim 7 wherein the calcium salt is calcium acetate.
11. The process of claim 7 wherein the calcium salt is dissolved in a solvent and the atorvastatin salt solution is contacted with the calcium salt by adding the calcium salt solution to the atorvastatin salt solution.
12. A process for preparing atorvastatin calcium Form V or hydrates thereof comprising the steps of
  - a) dissolving atorvastatin calcium in a solvent selected from the group consisting of tetrahydrofuran and hydroxylic solvents to form an atorvastatin calcium salt solution,
  - b) adding water to the atorvastatin calcium salt solution, and
  - c) isolating atorvastatin calcium Form V or hydrates thereof.
13. The process of claim 12 wherein the solvent is methanol.
14. The process of claim 12 wherein the solvent is ethanol.
15. The process of claim 12 wherein the solvent is tetrahydrofuran.
16. A pharmaceutical composition comprising a therapeutic amount of atorvastatin Form V or hydrates thereof of claim 1.

1/2



2/2



# INTERNATIONAL SEARCH REPORT

International application No.

PCT/US00/31555

## A. CLASSIFICATION OF SUBJECT MATTER

IPC(7) : C07D 207/335; A61K 31/40

US CL : 548/537; 514/423

According to International Patent Classification (IPC) or to both national classification and IPC

## B. FIELDS SEARCHED

Minimum documentation searched (classification system followed by classification symbols)

U.S. : 548/537; 514/423

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)

CAS ONLINE

## C. DOCUMENTS CONSIDERED TO BE RELEVANT

Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
A	US 5,969,156 A (BRIGGS et al) 19 October 1999 (19.10.1999).	1-16
A,P	US 6,121,461 A (MCKENZIE et al) 19 September 2000 (19.09.2000).	1-16

☐ Further documents are listed in the continuation of Box C.

☐ See patent family annex.

* Special categories of cited documents:	"T"
"A" document defining the general state of the art which is not considered to be of particular relevance	later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
"E" earlier application or patent published on or after the international filing date	"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)	"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art
"O" document referring to an oral disclosure, use, exhibition or other means	"&" document member of the same patent family
"P" document published prior to the international filing date but later than the priority date claimed	

Date of the actual completion of the international search

05 January 2001 (05.01.2001)

Date of mailing of the international search report

18 April 2001

Name and mailing address of the ISA/US

Commissioner of Patents and Trademarks

Box PCT

Washington, D.C. 20231

Facsimile No. (703)305-3230

Authorized officer

Fiona T. Powers

Telephone No. 703-308-1235